Amendments to the Claims

This claim listing replaces all prior versions and listings of claims in the application.

1-13. (Cancelled).

14. (Currently amended) A method of accelerating the start of growth of quiescent follicles in non-menopausal women comprising administering to a patient in need thereof a medicament comprising a somatostatin antagonist analog. The method of elaim 13, wherein the somatostatin antagonist analogue includes having the peptides of general formula (III)

$$\begin{array}{c} A^{1}\text{-cyclo}\{D\text{-Cys-}A^{2}\text{-}D\text{-Trp-}A^{3}\text{-}A^{4}\text{-Cys}\}\text{-}A^{5}\text{-}Y^{1}\\ (III) \end{array}$$

in which:

A1 is an optionally substituted aromatic α-amino acid;

 \boldsymbol{A}^2 is an optionally substituted aromatic $\alpha\text{-amino}$ acid;

A3 is Dab, Dap, Lys, or Orn;

 A^4 is β -Hydroxyvaline, Ser, Hser, or Thr;

 $\ensuremath{A^5}$ is an optionally substituted aromatic D- or L- $\alpha\mbox{-amino}$ acid; and

 Y^1 is OROH, NH_2 or NHR^1 , R^1 is (C_{1-6}) alkyl;

each aromatic α -amino acid being optionally substituted with one or more substituents independently includes selected from a halogen atom, NO₂, OH, CN, (C₁₋₆) alkyl, (C₂₋₆) alkynyl, (C₁₋₆) alkynyl, Q-Bzl or NR⁹R¹⁰, wherein R⁹ and R¹⁰ are each independently H, O, or (C₁₋₆) alkyl; and

each nitrogen atom with \underline{of} a peptide amide bond and the amino group of A^1 are optionally substituted with a methyl group, with the proviso that there is at least one said methyl group in a peptide of general formula (III);

the pharmaceutically acceptable salts or protected forms of said peptides , or combinations thereof.

- 15. (Cancelled).
- 16-18. (Cancelled).
- 19. (Currently amended) The method of claim 14, wherein the somatostatin antagonist analogue includes the peptide of formula (III), in which A^I is Cpa, A^2 is Pal, A^3 is Lys, A^4 is Thr, and A^5 is Nal.
- 20. (Currently amended) The method of claim 19, wherein the somatostatin antagonist analogue includes is Cpa e(DCys 3 Pal DTrp NMeLys Thr Cys) 2 Nal NH₂ Cpa-cyclo(DCys-3-Pal-DTrp-NMeLys-Thr-Cys)-2-Nal-NH₂.
- 21. (Currently amended) The method of claim 14, wherein the somatostatin antagonist analogue is: includes the peptide of formula (III), in which A[‡] is Cpa, A[‡] is 4Pal, A[‡] is Lys, A[‡] is Thr, and A[‡] is 2Nal

22. (New) A method of accelerating the start of growth of quiescent follicles in non-menopausal women comprising administering to a patient in need thereof a medicament comprising a somatostatin antagonist analog comprising:

the following peptides:

- Cpa-cyclo[D-Cys- Pal-D- Trp-N-Me-Lys- Thr-Cys]-D-Trp-NH₂;
- Cpa-cyclo[D-Cys-Tyr-D-Trp- N-Me-Lys-Thr-Cys]-Nal-NH₂;
- Cpa-cyclo[D-Cys-Pal-D- Trp- N-Me-Lys-Thr-Cys]- Nal-NH₂;
 the peptide acetyl-D-His-D-Phe-D-Ile-D-Arg-D-Trp-D-Phe-NH₂ (code name AC-178,335);

the octapeptide of the following structure (code name ODN-8);

the peptide Cpa-cyclo[D-Cys-Pal-D-Trp-Lys-Val-Cys]Cpa-amide (code name SB-710411);

the peptide of the following structure (code name BIM-23056);

BIM 23056

the compound of the following structure (code name BN-81674);

BN-81674

the compound of the following structure (code name SRA-880);

SRA-880

or their pharmaceutically acceptable salts or protected forms, or combinations thereof.